AMENDMENTS TO THE CLAIMS:

- 1. (Currently amended) A process of using a highly efficient catalyst-solvent system of dilute solutions of a base in water to effect a base catalyzed depolymerization (BCD) reaction for converting a biomass into a blending component for a petroleum-derived fuel comprising:
- (a) extracting a lignin-containing fraction in a reaction medium of water from the biomass to provide a lignin feed material;
- (b) base-catalyzed depolymerizing the lignin feed material in the presence of water and a low base concentration of about 2 to about 5 weight % in the absence of alcohol to provide a first composition comprising a depolymerized lignin conversion to ether solubles of between about 73 to about 74.5 weight percent; and
- hydrodeoxygenation and hydrocracking using a mixture of a co-catalyst system of a sulfided MMo/γ-Al₂O₃ for hydrodeoxygenation and a sulfided MMo/SiO₂-Al₂SO₃-zeolite for hydrocracking, where M=Group VI-VIII transition metal promoters of Ru, Co, Re, Cr, Fe, Pt and combinations thereof to provide a second composition comprising an aromatic hydrocarbon comprising C₇-C₁₀ alkylbenzenes of a purity of >90 weight%, for use as a high octane blending component for a-petroleum or lower grade petroleum-derived fuel gasoline.
- 2. (Canceled)
- 3. (Canceled)
- 4. (Original) The process of claim 1 wherein the blending composition is further defined as having a blending octane number of about 110 or higher.
- 5. (Canceled)
- 6. (Canceled)
- 7. (Original) The process of claim 1 wherein the biomass is a lignocellulose biomass.
- 8. (Currently Amended) The process of claim 1 wherein the second composition is further defined as comprising about 5% to 40% alkylated naphthenes.
- 9. (Currently Amended) The process of claim 8 wherein the second composition comprises about 5% to 30% alkylated naphthenes.
- 10. (Previously Amended) The process of claim 1 wherein the second composition is further

defined as comprising from about 75% to about 95% C_7 to C_{10} alkylbenzenes.

- 11. (Original) The process of claim 1 wherein the biomass is further defined as comprising from about 5% to about 70% lignin.
- 12. (Original) The process of claim 11 wherein the biomass comprises about 50% lignin.
- 13. (Original) The process of claim 1 or 12 wherein the biomass is further defined as comprising a Kraft lignin, organosolve lignin, a lignin derived from wood processing, a lignin as an ethanol process by-product, a lignin from a by-product of pulp and paper processing, or a combination thereof.
- 14. (Original) The process of claim 1 wherein the aqueous solvent comprises a dilute alkali hydroxide solution.
- 15. (Original) The process of claim 14 wherein the alkali hydroxide solution is about 0.5% to about 10% wt of an alkali hydroxide.
- 16. (Original) The process of claim 15 wherein the alkali hydroxide solution is about 0.5% to about 3% by wt NaOH.
- 17. (Original) The process of claim 1 wherein the depolymerization is carried out in the presence of a CsX-type zeolite as a superbase catalyst.
- 18. (Original) The process of claim 17 wherein the solid CsX-type zeolite superbase catalyst is used together with a 0.5% 5% alkali hydroxide solution as a co-catalyst system.
- 19. (Previously Amended) The process of claim 1 wherein the base-catalyzed depolymerization is further defined as occurring at an operational temperature of from about 300°C to about 340°C.
- 20. (Original) The process of claim 19 wherein the base-catalyzed depolymerization operation temperature is from about 310°C to about 330°C.
- 21. (Previously Amended) The process of claim 1 wherein the lignin-containing feed material has a liquid hourly space velocity of from 0.5h⁻¹ to about 10h⁻¹.
- 22. (Original) The process of claim 21 wherein the liquid hourly space velocity is from about 2.0h⁻¹ to about 9.0h⁻¹.
- 23. (Canceled)
- 24. (Canceled)
- 25. (Currently Amended) The process of claim 24 1 wherein the ratio of MMo/γ- Al₂O₃

catalyst: sulfided MMo/SiO2-Al2O3 zeolite catalyst is from 1:1 to 9:1.

- 26. (Cancelled)
- 27. (Original) The process of claim 1 wherein the hydroprocessing is further defined as comprising a hydrogen pressure of from about 500 psig to about 1800 psig.
- 28. (Original) The process of claim 27 wherein the hydrogen pressure is from about 500psig to about 1000 psig.
- 29. (Original) The process of claim 1 wherein the hydroprocessing is further defined as comprising a reaction temperature of from about 360°C to about 390°C.
- 30. (Original) The process of claim 29 wherein the reaction temperature is from about 380°C to about 390°C.
- 31. (Canceled)
- 32. (Canceled)
- 33. (Withdrawn) A biomass-derived blending component for a petroleum or petroleum-derived fuel comprising about 70% to about 95% C₇ to C₁₀ alkylbenzene.
- 34. (Withdrawn) The biomass-derived blending component of claim 33 further defined as comprising about 5% to about 30% alkylated naphthenes.
- 35. (Withdrawn) The biomass-derived blending component of claim 33 further defined as comprising about 5% to about 10% alkylated napthenes.
- 36. (Withdrawn) A method for enhancing the octane number of a petroleum or petroleum-derived fuel comprising: combining the biomass derived blending component of claim 33 with a petroleum or petroleum-derived fuel at a ratio of about 1:10 to about 1:4 by volume to provide a blended fuel, wherein the octane level of the petroleum blended fuel is enhanced about 1% to about 30% over the octane level of the petroleum or petroleum-derived based fuel without the blending component.
- 37. (Withdrawn) The method of claim 36 wherein the octane level of the blended fuel is enhanced about 30% over the petroleum or petroleum-derived fuel without the blending component.
- 38. (Withdrawn) The method of claim 36 wherein the petroleum or petroleum-derived fuel is gasoline.
- 39. (Currently Amended) A process of using a highly efficient catalyst-solvent system of

dilute solutions of a base in water to effect a base catalyzed depolymerization (BCD) reaction for converting a biomass into a blending component comprising a monocyclic aromatic hydrocarbon-rich composition for petroleum-derived fuels comprising:

- a) dispersing a lignin-containing feedstock material in a reaction medium of water to provide a dispersed lignin composition;
- b) base-catalyzed depolymerizing the dispersed lignin composition in the presence of water and a low base concentration of about 2 to about 5 weight percent in the absence of alcohol to provide a depolymerized lignin conversion to ether solubles of between about 73 to about 74.5 weight percent; and
- c) hydroprocessing the depolymerized lignin product by sequential or simultaneous hydrodeoxygenation and hydrocracking using a mixture of a co-catalyst system of sulfided MMo/γ-Al₂O₃ for hydrodeoxygenation and a sulfided MMo/SiO₂-Al₂SO₃-zeolite for hydrocracking, where M=Group VI-VIII transition metal promoters of Ru, Co, Re, Cr, Fe, Pt and combinations thereof to produce a high octane blending component comprising C₇ to C₁₀ alkylbenzenes of a purity >90 weight %.
- 40. (Canceled)
- 41. (Original) The process of claim 39 wherein the blending component comprises a blending octane number of about 95-150.
- 42. (Original) The process of claim 39 wherein the blending component is further defined as having a blending octane number of about 110.
- 43. (Canceled)
- 44 (Currently Amended) A process for producing BTX (benzene, toluene and xylenes) comprising:
 - a) extracting lignin-containing material from a biomass:
 - b) dispersing the lignin-containing material in a reaction medium of water;
- c) subjecting the dispersed lignin-containing material to a dilute solution of alkali hydroxide of about 2 to about 5 weight % in water in the absence of alcohol to produce a first composition comprising a depolymerized lignin conversion to ether solubles of between about 73 to about 74.5 weight percent; and
 - d) hydroprocessing the depolymenzed lignin by sequential or simultaneous

hydrodeoxygenation and hydrocracking using a mixture of a co-catalyst system of a sulfided MMo/γ-Al₂O₃ for hydrodeoxygenation and a sulfided MMo/SiO₂-Al₂SO₃-zeolite, where M=Group VI-VIII transition metal promoters of Ru. Co. Re. Cr. Fe. Pt and combinations thereof to provide a second composition comprising benzene, toluene, and xylenes, wherein the benzene, toluene, and xylenes for use as intermediates in the production of other organic chemicals.

- 45. (Currently Amended) A process of using a highly efficient catalyst-solvent system of dilute solutions of a base in water to effect a base catalyzed depolymerization (BCD) reaction for converting a biomass into a blending composition for a petroleum-derived fuel comprising:
 - a) extracting lignin-containing material from a biomass;
 - b) dispersing the lignin-containing material in a reaction medium of water;
- c) subjecting the dispersed lignin-containing material to a alkali hydroxide solution of about 2 to about 5 weight % alkali hydroxide to produce a first composition comprising a depolymerized lignin conversion to ether solubles of between about 73 to about 74.5 weight percent;
- d) hydrodeoxygenating and hydrocracking the first composition using a mixture of a co-catalyst system of a sulfided MMo/γ-Al₂O₃ for hydrodeoxygenation and a sulfided MMo/SiO₂-Al₂SO₃-zeolite, where M=Group VI-VIII transition metal promoters of Ru, Co, Re, Cr, Fe, Pt and combinations thereof to produce a second composition comprising C₇ to C₁₀ alkylbenzenes, wherein the second composition is used as a blending component for a petroleum-derived fuel.
- 46. (Canceled)
- 47 (Original) The process of claim 45 wherein the alkali hydroxide solution is about 1% to about 10% wt of an alkali hydroxide.
- 48. (Original) The process of claim 47 wherein the alkali hydroxide solution is about 1% to about 3% by wt NaOH.
- 49. (Canceled)
- 50. (Original) The process of claim 47 wherein the second composition is further defined as comprising from about 75% to about 95% C₇ to C₁₀ alkylbenzenes.